Billing Code 4410-09-M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer Of Controlled Substances; Notice Of Application; Apertus Pharmaceuticals, LLC

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on March 27, 2012, Apertus Pharmaceuticals, LLC, 331 Consort Drive, St Louis, Missouri 63011, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Alfentanil (9737)	II
Remifentanil (9739)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The company plans to manufacture small quantities of the listed controlled substances to make reference standards which will be distributed to their customers.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substances, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than [Insert date 60 days from date of publication].

Joseph T. Rannazzisi
Deputy Assistant Administrator
Office of Diversion Control
Drug Enforcement Administration

Dated: June 4, 2012

[FR Doc. 2012-14165 Filed 06/11/2012 at 8:45 am;
Publication Date: 06/12/2012]